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APPLICATION NO.		FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
10/657,205		09/09/2003	Narutoshi Fukuzawa	242335US0	9140		
22850	7590	10/24/2006		EXAM	EXAMINER		
C. IRVIN I	MCCLE	LLAND	ANGEBRANNDT, MARTIN J				
OBLON, SF 1940 DUKE	•	ИССLELLAND, М Г	ART UNIT	PAPER NUMBER			
ALEXAND	RIA, VA	22314	1756				
					DATE MAILED: 10/24/2006		

Please find below and/or attached an Office communication concerning this application or proceeding.

		Applic	ation No.	Applicant(s)	P			
			7,205	FUKUZAWA ET AL.				
Offic	e Action Summary	Exami	ner	Art Unit				
		Martin	J. Angebranndt	1756				
The MAI Period for Reply	LING DATE of this communication	ation appears on	the cover sheet w	ith the correspondence add	ress			
	O CTATUTODY DEDICO FO		TO EVOIDE AL	AONTHION OF THIRTY (20)	NDAYC			
WHICHEVER I - Extensions of time after SIX (6) MONT - If NO period for rep - Failure to reply with Any reply received	D STATUTORY PERIOD FOI S LONGER, FROM THE MAI may be available under the provisions of FHS from the mailing date of this commun sly is specified above, the maximum statu- in the set or extended period for reply will by the Office later than three months after adjustment. See 37 CFR 1.704(b).	ILING DATE OF 37 CFR 1.136(a). In no ication. tory period will apply an II, by statute, cause the	THIS COMMUNI be event, however, may a and will expire SIX (6) MOI application to become Al	CATION. reply be timely filed NTHS from the mailing date of this com BANDONED (35 U.S.C. § 133).				
Status								
1)⊠ Responsi	ive to communication(s) filed	on 09 August 20	006.					
2a) This action	·)⊠ This action i						
3) Since this	·							
closed in	accordance with the practice	under <i>Ex parte</i>	Quayle, 1935 C.). 11, 453 O.G. 213.				
Disposition of Cla	ims							
4)⊠ Claim(s)	1-3 and 5-16 is/are pending i	n the application						
•	above claim(s) is/are							
5) Claim(s)	is/are allowed.							
6)⊠ Claim(s)	1-3 and 5-16 is/are rejected.							
•	is/are objected to.		•					
8) Claim(s)	are subject to restriction	on and/or electio	n requirement.					
Application Paper	S							
9)∐ The speci	fication is objected to by the	Examiner.						
10)☐ The drawi	ing(s) filed on is/are: a	a) accepted or	b) objected to	by the Examiner.				
Applicant	may not request that any objection	on to the drawing(s) be held in abeya	nce. See 37 CFR 1.85(a).				
•	ent drawing sheet(s) including the							
11)☐ The oath	or declaration is objected to b	by the Examiner.	Note the attache	d Office Action or form PTC)-152.			
Priority under 35 l	J.S.C. § 119				•			
12) Acknowle	dgment is made of a claim fo	r foreign priority	under 35 U.S.C.	§ 119(a)-(d) or (f).				
a)∐ All b)	Some * c) None of:							
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Attachment(s)				·				
1) Notice of Referen	ices Cited (PTO-892)		4) Interview	Summary (PTO-413)				
2) D Notice of Draftspo	erson's Patent Drawing Review (PTC	D-948)	Paper No	(s)/Mail Date				
3) Information Discle Paper No(s)/Mail	osure Statement(s) (PTO/SB/08) Date		6) Other:	Informal Patent Application				

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1. The response of the applicant has been read and given careful consideration. Responses to the arguments relating to the rejection are presented after the first rejection to which they are directed. The amendment to the specification is accepted. Rejections of the previous office action, not repeated below are withdrawn based upon the amendment and acc0ompaniying arguments.

- 2. The following is a quotation of the second paragraph of 35 U.S.C. 112: .
 - The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.
- 3. Claim 5 is rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

This claim should state that the two heterocyclic rings are both benzoxazole. The current claims mostly repeat limitations in claim 1 and adds only that the heterocyclic groups are the same. Claims 1 limits this case to where they are both benzoxazole.

- 4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 5. Claims 1-3,5 and 8-16 are rejected under 35 U.S.C. 103(a) as being unpatentable over JP 10-188339, in view of Sano et al. JP 06-044608.

JP 10-188339 teaches the use of cyanine dyes including indolene, benzodiazole and benzooxazole dyes embraced by formula I, where A moeties form a phenyl or naphthyl ring, the

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counterions can be halogen and the like. The N substituents can be C₁₋₆ alkyl. [0007,0016,0033-0034]. Formula (3) is a benzodiazole trimethine dye, formula (4) is a indolenic trimethine dye, and formula (5) is a benzooxazole trimethine dye, where the N substitutents are methyl or ethyl and the counterion is iodine. [0043]. In the examples 1-4, a polycarbonate substrate is coated with the dye layer, a gold reflective layer and a UV curing layer applied [0041]. In example 4, dye 4, (of formula 5) is used. [0042-0043]. The use of inorganic layers as interference or solvent resistance layers is disclosed [0036-0037].

Sano et al. JP 06-044608 teaches a dielectric layer (31) formed on the recording layer (2) and another dielectric layer (32) between the reflective layer (4) and the UV curable layer (5). The dielectric layers are inorganic films [0011] and increases the adhesion between the layers of the media and prevents damage due to high humidity/temperature [0004-0006,0014]. The recording layers can be cyanine dyes [0009].

It would have been obvious to one skilled in the art to modify example 4 of JP 10-188339 by adding dielectric layers on one of both sides of the reflective layer as taught by Sano et al. JP 06-044608 with a reasonable expectation of increasing the adhesion between the layers and preventing damage to the medium based upon humidity/temperature based upon the direction to the use of inorganic layers within JP 10-188339 at [0036-0037] and the discussion of cyanine dyes in both of the references. Further, it would have been obvious to one skilled in the art to modify the resulting example by replacing the N alkyl groups with butyl based upon the disclosure of equivalence for the alkyl groups at [0017].

Claim 1 does not require that the light transmitting layer be directly in contact with the dielectric layer nor does it state that the medium is capable of being recorded and/or reproduced

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thereon with laser light of 390-420 nm incident upon the light transmitting layer surface. The claims merely set forth a material limitation for the light transmitting layer and does not speak to the remainder of the medium. With respect to the limitations regarding the refractive index, these are inherent properties of the dyes and while the media must have a reasonable expectation of functioning with a laser within the 380-420 nm wavelength range, the claims to the article embrace the media irrespective of what wavelength they are used with.

The applicant argues the test data. The examiner notes that the dye of formula (5) in section [0043] of JP 10-188339is identical to dye AA-1 Of the instant specification. Further the articles claims are not limited to use with a particular laser or wavelength.

6. Claims 1-3,5,6 and 8-16 are rejected under 35 U.S.C. 103(a) as being unpatentable over JP 10-188339, in view of Sano et al. JP 06-044608, further in view of Sato et al. '839.

Sato et al. '839 teach the increase in the solubility of cyanine dyes when the N substituents are different from one another (abstract and 2/4-15). The addition of stabilizers (quenchers) to recording layers is disclosed as enhancing the stability of recording layer (21/57-23/35).

In addition to the basis set forth above, it would have been obvious to modify the media rendered obvious by JP 10-188339, in view of Sano et al. JP 06-044608 by adding a stabilizer (quencher) to improve the stability of the cyanine dyes based recording layer as taught by Sato et al. '839. Further, it would have been obvious to one skilled in the art to modify the resulting example by using a butyl moiety as the N substitutent on the benzoxazole moiety and replacing the other benzoxazole moiety with either an indolene or benzodiazole moiety with a methyl

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and/or ethyl moieties as the N substitutent based upon the direction in formula I of JP 10-188339 and the direction within Sato et al. '839 that this increases the solubility of the dyes.

7. Claims 1-3,5 and 7-16 are rejected under 35 U.S.C. 103(a) as being unpatentable over either of Berneth et al. '807, in view of JP 10-188339.

Berneth et al. '807 teach optical recording media read from the side opposite the substrate which increases the resolution (near field recording) [0010-0011,0018]. The use of cyanine dyes in the recording layer is disclosed [0024]. The recording takes place between 360 and 460 nm with a high NA. [0018]. Embodiment 7 uses a trimethine cyanine dye, have the recording layer on the substrate, a 40 nm silicon dioxide dielectric layer (12) applied to that and a protective layer adhered via an adhesive layer and was recorded on using a 405 nm laser from the light transmitting layer side [0031-0038,0040-0044,page 7]. The dielectric layer protects the recording layer.

It would have been obvious to one of ordinary skill in the art to modify the examples of Berneth et al. '807 by using other trimethine cyanine dyes embraced by formula I within JP 10-18, including those exemplified or rendered obvious by JP 10-188339 as discussed above with a reasonable expectation of forming a medium useful with a 405 nm laser as shown in example 7 Further, it would have been obvious to record on the resulting media using lasers emitting in the 380-425 nm range, particularly the 405 nm laser exemplified.

In addition to the basis above, the examiner notes that the dye used in example 7 differs from those tested by the applicant in table 1 of the instant specification. The Dye used is a benzothiozole analog of dye AC-1. If the applicant can show a difference when these are used ina declaration, the rejection made be obviated. The examiner notes that there is no showing for

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Dye AC-1 or any of the asymmetric dyes exemplified by the applicant in table 1. The applicant argues that a wide range of dyes are disclosed in Berneth et al. '807, but neglects that there is a specific example which presents an opportunity for comparative data.

8. Claims 1-3 and 5-16 are rejected under 35 U.S.C. 103(a) as being unpatentable over either of Berneth et al. '807, in view of JP 10-188339, further in view of Sato et al. '839 and Sabi et al. EP 1103962.

Sabi et al. EP 1103962 teaches a topside accessed optical recording medium having the structure shown in the figures. A dielectric layer is provided between the recording layer and the protective layer to prevent the dissolution/damage of the recording layer during coating of the UV curing resin protective layer [0011,0042]. Adjustment of the thickness of the dielectric layer can also be used to optimize the signal amplitude [0064].

In addition to the basis set forth above, it would have been obvious to modify the media rendered obvious by the combination of Berneth et al. '807 and JP 10-188339 by adding a stabilizer (quencher) to improve the stability of the cyanine dyes based recording layer as taught by Sato et al. '839. Further, it would have been obvious to one skilled in the art to modify the resulting example by using a butyl moiety as the N substitutent on the benzoxazole moiety and replacing the other benzoxazole moiety with either an indolene or benzodiazole moiety with a methyl and/or ethyl moieties as the N substitutent based upon the direction in formula I of JP 10-188339 and the direction within Sato et al. '839 that this increases the solubility of the dyes and further the evidence in Sabi et al. EP 1103962 describes the damage/degradation of the recording layer by direct application of the protective layer upon I and that the use of a dielectric interlayer solves this problem.

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 The prior art made of record and not relied upon is considered pertinent to applicant's disclosure.

Kamezaki et al. JP 03-146393 teach the use of cyanine dyes in the examples, including the benzooxazole trimethine dye (1) in the lower right column on page 3. The N substitutents are both ethyl and the counter ion iodine.

Tominaga et al. JP 10-168450 teach the use of cyanine dyes including indolene, benzooxazole dyes embraced by formula 19, where the counterions can be I- and the like. The N substituents can be C_{1-10} alkyl. [0032-0034]. A trimethine indolene dyes is shown in formula 20 [0041] and an oxazole dyes is shown in formula 22, where the N substitutent is propyl [0048-0050].

JP 60-204395 teaches the provision of a silicon dioxide layer on both sides of the cyanine dye based recording layer.

10. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Martin J. Angebranndt whose telephone number is 571-272-1378.

The examiner can normally be reached on Monday-Thursday and alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mark Huff can be reached on 571-272-1385. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Martin J Angebranndt Primary Examiner

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10/20/2006